




A global assessment of plant–mite mutualism and its ecological drivers

Andrew Myers^{a,b}, Bruce Martin^{a,c}, Jenna Yonenaga^d, Anurag A. Agrawal^{e,1} , and Marjorie G. Weber^{a,1}

Affiliations are included on p. 9.

Contributed by Anurag A. Agrawal; received June 6, 2023; accepted September 26, 2024; reviewed by Megan E. Frederickson and Brian C. O'Meara

Mutualisms are mediated by adaptive traits of interacting organisms and play a central role in the ecology and evolution of species. Thousands of plant species possess tiny structures called “domatia” that house mites which protect plants from pests, yet these traits remain woefully understudied. Here, we release a worldwide database of species with mite domatia and provide an evaluation of the phylogenetic and geographic distribution of this mutualistic trait. With >2,500 additions based on digital herbarium scans and published reports, we increased the number of known species with domatia by 27% and, importantly, documented their absence in >4,000 species. We show that mite domatia likely evolved hundreds of times among flowering plants, occurring in an estimated ~10% of woody species representing over a quarter of all angiosperm families. Contrary to classic hypotheses about the evolutionary drivers of mutualism, we find that mite domatia evolved more frequently in temperate regions and in deciduous lineages; this pattern is concordant with a large-scale geographic transition from predominantly ant-based plant defense mutualisms in the tropics to mite-based defense mutualisms in temperate climates. Our data also reveal a pattern of evolutionary convergence in domatia morphology, with tuft-form domatia more likely to evolve in dry temperate habitats and pit domatia more likely to evolve in wet tropical environments. We have shown climate-associated drivers of mite domatia evolution, demonstrating their utility and power as an evolutionarily replicated system for the study of plant defense mutualisms.

acarodomatia | global plant database | mite domatia | indirect plant defense | mutualism

Mutualisms, or cooperative interactions between species, play a central role in the generation and maintenance of biodiversity (1). Plants commonly engage in mutualistic interactions for nutrient exchange, defense, and reproduction and have evolved a suite of remarkable adaptive traits for attracting and retaining mutualists. Because the primary function of these phenotypes is to mediate interactions with mutualists, they serve as excellent models for testing fundamental hypotheses at the intersection of ecology and evolutionary biology (2). For example, Dobzhansky's classic “Biotic Interactions Hypothesis” posits that the selective pressure of interspecific interactions is relatively stronger in tropical over temperate habitats, favoring investment in adaptations for mutualistic interactions such as animal pollination (3–6). Similarly, traits that facilitate mutualisms have been classically hypothesized to evolve more frequently where both opportunities for cheating and costs are low (4, 7), or where stress is high [e.g., “Stress Gradient Hypothesis” (8)], leading to testable, yet sometimes contradictory, predictions for geographic patterns in investment in mutualistic phenotypes at large scales.

One of the world's most ancient and widespread—yet globally understudied—plant mutualistic interactions is between plants and mites (9, 10). In plant–mite systems, plants possess mite domatia (also known as “acarodomatia” or “leaf domatia”)—small, covered chambers generally on the abaxial vein axils of leaves that provide shelter for beneficial mites (10–12). Unlike galls, mite domatia are present regardless of mite occupants (i.e., they are not induced by the presence of mites) (13). They generally take the form of small depressions in the leaf surface covered by either a dense layer of trichomes (referred to as “tuft” domatia), an open flap of laminar tissue (“pocket” domatia), or a closed flap of laminar tissue with a small pore (“pit” domatia). All three morphologies create semicovered chambers that are often occupied by the eggs, nymphs, and adults of predacious, fungivorous, and scavenging mites (typically Acariformes).

A substantial body of evidence reveals that mite domatia enhance natural enemy abundances on leaves (13–26). Mites protect plants by consuming small herbivores and fungal pathogens (13, 19), and in return, mites benefit from the shelter afforded by the domatia through mechanisms such as protection from abiotic stresses and escape from enemies (14–18, 27). By housing a “standing army” (28) of mite bodyguards, mite domatia provide

Significance

Mite domatia are small structures on leaves common in woody flowering plants. These structures serve as domiciles for tiny mites that typically protect plants by consuming fungal and arthropod pests. We estimate that over 14,000 plant species have these protective structures and ask where they are found across the globe. Mite domatia evolved hundreds of times, suggesting a repeated adaptive response to harness protection from beneficial mites. We found a broad scale association between mite domatia and temperate climates, reflecting a trade-off in the proportion of plants cooperating with ants versus mites moving from the tropics toward the poles. The open-access database we release with this study will serve as a foundation for future studies of this exceedingly common mutualism.

Author contributions: M.G.W. designed and funded the original research study; A.M., B.M., J.Y., A.A.A. and M.G.W. developed the study and performed research; A.M., J.Y. and M.G.W. collected data; A.M., B.M., and M.G.W. analyzed data; and A.M., B.M., A.A.A., and M.G.W. wrote the paper.

Reviewers: M.E.F., University of Toronto; and B.C.O., University of Tennessee.

The authors declare no competing interest.

Copyright © 2024 the Author(s). Published by PNAS. This article is distributed under [Creative Commons Attribution-NonCommercial-NoDerivatives License 4.0 \(CC BY-NC-ND\)](https://creativecommons.org/licenses/by-nc-nd/4.0/).

¹To whom correspondence may be addressed. Email: aa337@cornell.edu or webermg@umich.edu.

This article contains supporting information online at <https://www.pnas.org/lookup/suppl/doi:10.1073/pnas.2309475121/-/DCSupplemental>.

Published November 19, 2024.

Table 1. Total numbers and percentages of plant species, genera, and angiosperm families with mite domatia

Mite domatia presence	Number of species (% total spp. in list)	Number of genera represented	Number of families represented	Mite domatia type	Number of species (% of mite domatia spp.)
Present	2,514 (39%)	565	105	Tuft	674 (27%)
				Pit	496 (20%)
				Pocket	552 (22%)
				Variable	143 (6%)
				Unknown	649 (26%)
Absent				4,014 (61%)	641
List total	6,528 (100%)			1,070	141

For species with mite domatia present, domatia morphological type percentages 149 are tallied.

powerful protection to plants, reducing leaf damage in many woody species living in various ecosystems across the globe. Surveys of plant communities revealed that species with mite domatia make up to 58% of native tree species in North American deciduous forests (20), 42% of woody species in deciduous coastal forests of Korea (21), and 31% of woody species in southern Islands of New Zealand (22). Mite domatia also occur on economically important crops, including grapes, cherries, peppers, maples, oaks, and coffee (11). Despite their ubiquity and economic significance, we still know strikingly little about mite domatia, including the fundamental patterns of their phylogenetic and geographic distribution.

There are several hypotheses about specific factors that shape the phylogenetic and geographic distribution of mite domatia. On the one hand, the ubiquitous presence of beneficial mites as “aerial plankton” could result in relatively uniform selection across ecosystems for mite domatia as low-cost and reliable mutualistic phenotypes, suggesting that domatia should be evenly distributed across ecosystems. On the other hand, evolutionary transitions involving mite domatia, including gains, losses, and morphological changes appear common among closely related plant species (22, 23), and some plant communities have higher incidence of domatia, suggesting that specific conditions may favor their evolution. For example, mite domatia have been hypothesized to be most beneficial in dry, warm environments where they may better protect mites from desiccation (24, but see ref. 25), or in wet environments where higher herbivore and fungal loads impose stronger burdens on leaves (11, 15, 29). Similarly, some suggest mite domatia occur more frequently in temperate plant species due to increased abiotic stress and risk of mite-desiccation, predicting positive correlations between latitude and domatia occurrence (12). However, this suggestion has never been tested, and it contrasts with general hypotheses about the evolution of defense and mutualism, which predict traits like domatia to be evolutionarily associated with lower latitudes due to high diversity and abundance of herbivores, pathogens, and mutualists (3, 4, 6, 30, 31). Finally, interactions with other plant traits may influence selection on mite domatia. For example, longer-lived leaves are expected to be exposed to greater herbivory and thus experience selection for stronger defenses (32). Despite myriad hypotheses, the distribution of mite domatia across the plant tree of life was previously unknown, preventing phylogenetically controlled tests of these hypotheses on a broad scale (23, 33–35).

Here, we construct a global database of species with mite domatia and estimate the total number of species with these mutualistic morphological traits worldwide. We pair our database with a modern phylogeny of plants to assess the macroevolutionary distribution of mite domatia, patterns of phylogenetic convergence, and the extent to which species with mite domatia associate with

different geographic regions repeatedly across clades. Using phylogenetically controlled analyses, we test for evidence that previously hypothesized factors such as precipitation, temperature, latitude, and leaf lifespan evolutionarily correlate with the presence and morphology of mite domatia across the plant tree of life. Finally, given the well-known increase in ant defense of plants at lower latitudes (36) we coupled phylogenetic and geographic analyses to address whether patterns of plant–mite associations are concordant with or trade-off against patterns of plant associations with ants.

Results

One in 10 Woody Plants Is Estimated to Provide Housing for Protective Mites. We compiled species reported to have mite domatia in the literature (species descriptions, ecological studies, systematic surveys, etc.), revealing domatia in 2,514 species representing over 25% of angiosperm families (565 genera and 105 families, Table 1, *SI Appendix, Table S1*, and *Dataset S1*), increasing the known number of plants previously reported to have mite domatia (11) by 27% (685 new species, 148 new genera, and 13 new families). 4,014 species were well documented to lack domatia. Mite domatia were found almost exclusively on woody plants, with the few exceptions occurring almost entirely in cultivated species (*SI Appendix, Table S2*). Using a taxonomically informed Bayesian model, we estimated the total number of woody angiosperm species with mite domatia (beyond those documented in our list). Using a conservative model, we estimated 9.7 to 10.9% of woody angiosperms species possess mite domatia (median = 10.3%, roughly 14,200 to 16,800, based on 95% CI). These estimates suggest our current list of ~2,500 species represents roughly one sixth of total domatia-bearing diversity and that ~10% of woody angiosperms likely engage in a defense mutualism with mites worldwide.

Distantly Related Woody Plants across the Plant Phylogeny Have Convergent Evolution of Mite Domatia. To assess the macroevolutionary distribution of mite domatia and patterns of phylogenetic convergence, we placed our dataset on the GBOTB extended mega-tree of vascular plants (37). Species with mite domatia were widely distributed across the plant phylogeny, occurring in 64% of angiosperm orders (Fig. 1 and *SI Appendix, Fig. S2 and Table S1*). Although our scale of analysis and the rapid rate of domatia evolution inherently limits specific estimates of evolutionary transitions, it is clear that mite domatia evolved a striking number of times across the plant tree of life, with stochastic character mapping using the best fit model suggesting that domatia evolved 600 to 700 times and were lost 500 to

600 times among the species in our list. Evolutionary rates for domatia were heterogeneous among clades, with particular lineages displaying high capacity for domatia evolution compared to others (*SI Appendix, Fig. S1*, $\Delta AIC M1-M0 = 191.2$). Domatia-bearing species were most numerous among eudicots (present within 33% of eudicot families; Fig. 1 and *SI Appendix, Fig. S2*) and rare among monocotyledonous species (with only three species represented, likely due to parallel vein architecture limiting the evolution of vein-axil related structures such as domatia). Similarly, mite domatia are rare among the other noneudicot lineages, with the exception of the magnoliid group, which has prominent vein axils. Within Eudicots, four families together represented nearly half of all known mite domatia-bearing species (Rubiaceae 819 spp., 33% of all species in our list; Sapindaceae 124 spp., 5%; Combretaceae 122 spp., 5%; Melastomataceae 112 spp., 4 %; in total, these families represent ~11% of woody eudicots) (*SI Appendix, Fig. S2* and *Table S1*). Together, this represents a striking level of convergence, whereby lineages separated by hundreds of millions of years of evolution have

converged on woody lifestyles and evolved adaptations to house mutualistic mite defenders.

Ecological Predictors of Repeated Mite Domatia Evolution.

To test whether the highly convergent evolutionary history of mite domatia is consistent with hypotheses about the drivers of mutualism, we conducted phylogenetic logistic regressions between mite domatia (presence/absence and morphology) and a) geographic factors (latitude and elevation), b) climatic factors (temperature and precipitation), and c) plant phenotypes (phenology and leaf thickness). We evaluated the contemporary geographic distribution of mite domatia via occurrence data for each species using the Global Biodiversity Information Facility (GBIF) (38, 39) and acquired plant trait data from the TRY database (40). We collated over 1.1 million herbarium record coordinates for 5,544 species (85% of our list), with a mean number of 199 coordinates per species (min = 1, max = 7,896 coordinates). Plants with mite domatia were widely distributed

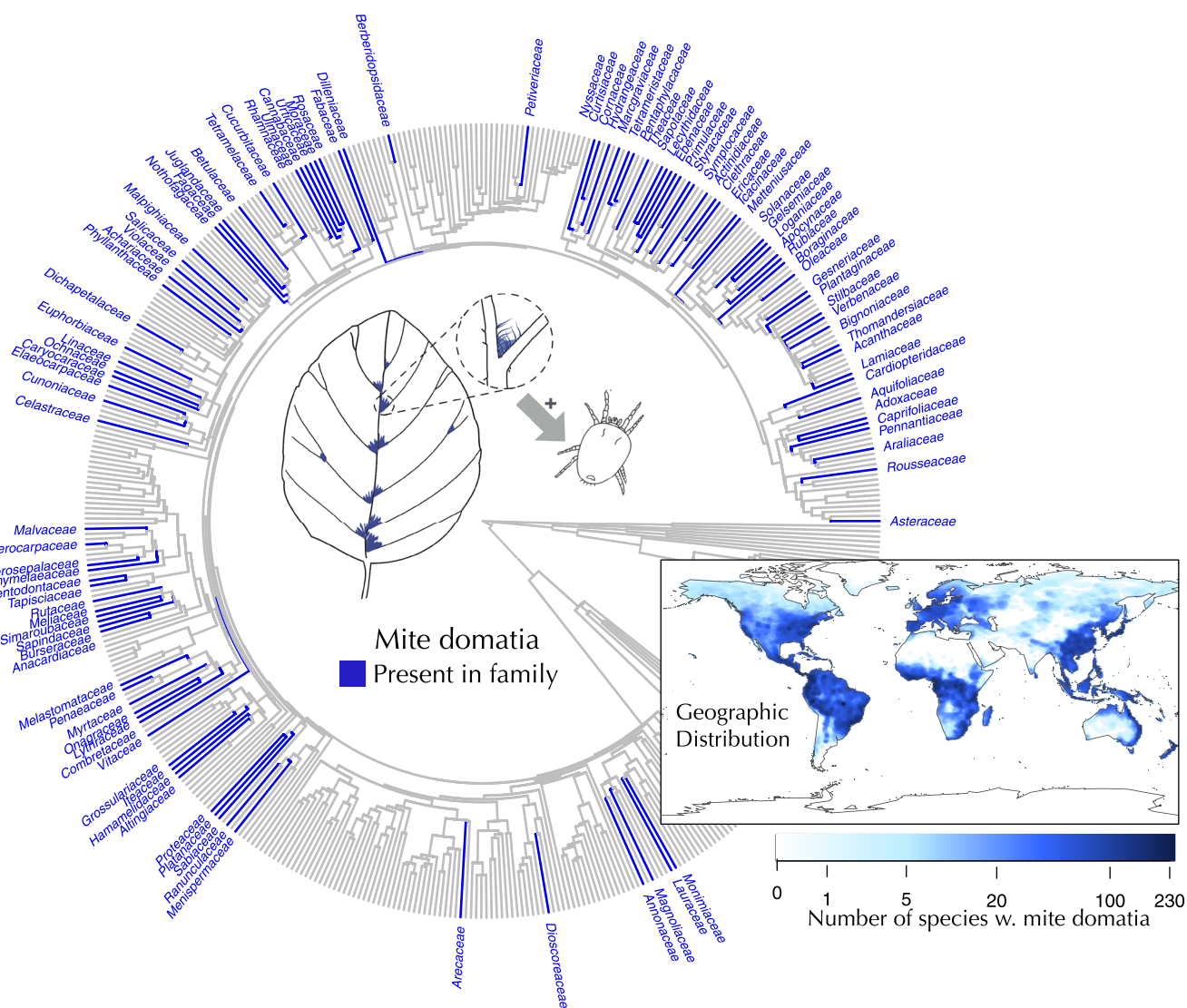


Fig. 1. Mite domatia are small, convergently evolved plant structures that mediate a mutualism between predacious and fungivorous mites and plants. Mite domatia generally occur in vein axils on the undersides of plant leaves. The structures provide shelter for beneficial mites which consume microbes and herbivores on the leaf surface. (Center) The phylogenetic distribution of mite domatia plotted on the GBOTB extended mega-tree of vascular plants. Each tip represents a plant family, with blue lineages denoting families that contain species with mite domatia. The area of the phylogeny covered by the map does not contain any instances of mite domatia and represents gymnosperms, ferns, ancient angiosperm lineages, and a portion of monocotyledonous families. (Bottom right) The geographic distribution of plants with mite domatia. Blue shading represents the number of species present with mite domatia present in an area.

Table 2. Estimated effect sizes of relationships between mite domatia occurrence and morphology predicted by climate and other plant traits



Predictor Variable (units of change)	Overall	Tuft	Pit	Pocket
Mean annual temperature (1 °C Increase)	-3%**	-5%*	4%	3%
Total annual precipitation (10 cm increase)	-3%**	-3%**	6%***	-3%*
Latitude (1° poleward)	2%**	2%***	-2%	1%
Elevation (100 m increase)	2%*	-3%*	2%	2%
SLA (10% increase)	5%***	2%	-6%**	3%
Mixed phenology (compared to deciduous)	-9%	-21%**	119%**	-23%
Evergreen phenology (compared to deciduous)	-45%***	-53%**	354%***	-24%

Effect sizes are reported as percent changes in odds of having mite domatia or a particular domatia morphology given a change in the predictor variable in the units reported under Predictor Variables. The last two rows relate to discrete predictor variables and effect sizes signify a change in odds across groups (e.g., for the last row, mite domatia are 45% less likely to occur in evergreen lineages compared to deciduous lineages). Significant relationships are bold-faced, with ***, **, and * indicating $P < 0.001$, 0.01, and 0.05, respectively. All P -values are estimated from simulated null distributions of slope values for each predictor separately (see *SI Appendix, Fig. S5* for correlations between predictors) and adjusted for multiple testing based on the Benjamini-Hochberg procedure. Samples sizes for analyses are listed in *SI Appendix, Table S3*. Domatia illustrations by John Megahan.

across the globe (Fig. 1 and *SI Appendix, Fig. S3*). The probability of observing mite domatia was positively associated with latitude and elevation and negatively associated with mean annual temperature and rainfall (Table 2 and *SI Appendix, Fig. S4* and Table S2), reflecting the highly repeated occurrence of mite domatia in distantly related lineages of plants in colder, drier environments (i.e., convergence in temperate ecosystems). We found this association despite large clades of closely related species with domatia in our dataset occurring in the tropics (for example, the tropical family Rubiaceae, representing >33% of our species). Mite domatia were also positively associated with deciduous leaves and higher specific leaf area (SLA) (i.e., thinner leaves with greater surface area-to-mass ratios) across the phylogeny (Table 2 and *SI Appendix, Fig. S4*).

Our finding that the presence of mite domatia is positively correlated with latitude is in stark contrast to patterns of evolution found for other plant defense mutualism traits associated with ants [myrmecodomatia and extrafloral nectaries (EFNs)], which are correlated with tropical, lower latitudes (41). Thus, as one moves away from the equator toward the poles, mites may predictably replace ants as plant bodyguards. To test for evolutionary and geographic patterns consistent with this hypothesis, we created a combined dataset of plants scored as having mite defense (mite domatia) vs. ant defense (either EFNs or ant domatia) phenotypes. Tests of correlated evolution revealed a strong negative evolutionary association between mite and ant defense across the phylogeny [corHMM Δ AICc (corrected Akaike information criterion) independent – correlated model = 517.32; phylogenetic logistic regression: $P < 0.001$; *SI Appendix, Fig. S6*]. As predicted, there was a strong latitudinal structure of turnover between mite and ant defense across plants (Fig. 2), with the proportion of species with mite, rather than ant, defense increasing exponentially moving away from the tropics toward the poles ($R^2 = 0.37$, $\beta = 0.01$, $P = 0.0003$).

Large-Scale Latitudinal Convergence in Domatia Morphology: Tufts, Pockets, and Pits. Incorporating morphological data into phylogenetic analyses revealed widespread convergence across the

three main morphological forms of mite domatia (tufts, pockets, and pits). Of the species with mite domatia, 68% could be confidently categorized into one of the three primary morphotypes using photographs, herbarium records, or written descriptions of the morphology. Morphotypes were similarly represented across the dataset: 39% (674 species) had tuft domatia, 32% (552 species) had pocket domatia, and 29% (496 species) had pit domatia (Table 1). Mite domatia were variably present or had multiple morphotypes reported in 429 species, reflecting intraspecific variation (Table 1).

Of the families represented in our list, 25/121 families had representatives of all three forms, a clear signature that the forms repeatedly evolved across distantly related groups (Fig. 3 and *SI Appendix, Fig. S2*). Analyses broken down by morphotype (tuft, pocket, or pit) reveal divergent associations with geographic and climatic variables (Table 2 and *SI Appendix, Fig. S4*). Most notably, tuft domatia were associated with higher latitudes across the phylogeny, while pit domatia were more prevalent in lower latitudes. Pit domatia repeatedly occurred in association with higher precipitation and marginally with higher temperatures, while tuft domatia showed negative associations with precipitation and temperature (Table 2 and *SI Appendix, Fig. S4*). With the exception of a negative correlation with precipitation, pocket domatia showed no significant relationships with climatic or geographic variables, reflecting their relatively even distribution across biomes. Together, these results reflect a scenario of repeated geographic convergence in domatia forms to different climates: pit domatia convergence in tropical (warm, wet) habitats and tuft domatia convergence in temperate (cold, dry) zones. Like climatic predictors, we found differences among domatia types had distinct associations with leaf traits, reflecting patterns of convergence across mite domatia morphology in certain types of plants (Table 2 and *SI Appendix, Fig. S4*). Pit domatia were repeatedly associated with thicker, evergreen leaves, whereas tuft domatia were associated with deciduous leaves and did not show any association with SLA. Pocket domatia were relatively evenly distributed among plants exhibiting deciduous, evergreen, and variable phenology types.

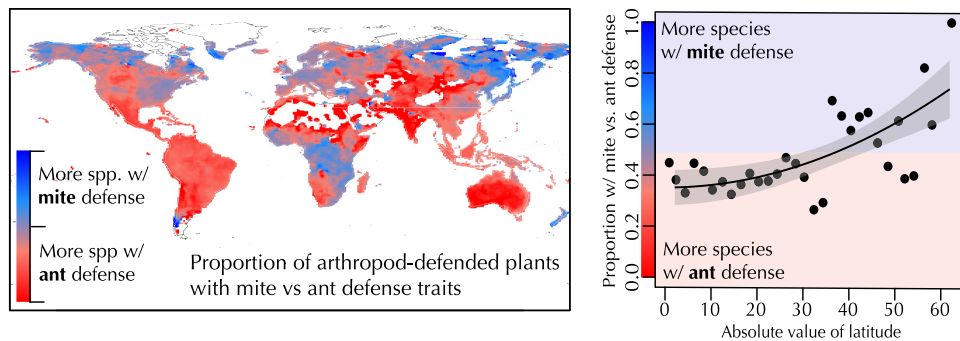


Fig. 2. The geographic turnover in investment in mite vs. ant defense. (Left) The relative proportion of known plants with mite domatia relative to plants with ant defense traits (EFNs vs. ant domatia) across the globe. (Right) The proportion of species with mite domatia increases with latitude. Points represent the number of species with mite domatia divided by the total number of species with either ant or mite mutualism traits in 3° bands of latitude, with 1 representing all of the plant species with mutualistic defense traits having mite domatia and 0 representing all of the species with mutually defended traits having either EFNs or ant domatia. Species were assigned to bins based on the median latitude of their range.

Discussion

We provide a large-scale assessment of the phylogenetic and geographic distribution of mite domatia, one of earth's most widespread convergently evolved plant defense mutualism traits. Our study revealed three main findings. First, mite domatia are widely distributed in plant species across the plant tree of life, estimated to occur in 10% of woody angiosperm species worldwide, suggesting a highly repeated adaptive response in plants to harness protection from beneficial mites. Second, phylogenetic tests revealed a widespread evolutionary correlation between mite domatia and temperate climatic regions with relatively lower temperatures and rainfall, reflecting a broad-scale turnover from ant to mite defense as one moves from the tropics to the poles. Third, the three common forms of mite domatia (tufts, pockets, and pits) showed distinct geographic distributions, reflecting a large-scale convergence across geographic regions and leaf phenologies (deciduous/evergreen), indicating differential adaptation to particular environments and plant forms.

Evolutionary rates for the origin or loss of domatia were heterogeneous among plant clades, with some lineages displaying high capacity for domatia evolution compared to others. However, our large dataset, phylogenetically informed models, and bootstrapping approach allowed us to tease apart taxonomically driven and geographical-sampling biases to identify evolutionarily replicated associations between mite domatia and hypothesized predictor variables. Our approach reveals that domatia are positively associated with latitude and elevation and negatively associated with mean annual temperature, precipitation, and evergreenness across the plant phylogeny (Table 2 and *SI Appendix, Fig. S4*). Our analyses provide a test of the hypothesis that mite defense is more common in temperate, rather than tropical regions, a pattern that conflicts with the prominent and classic Biotic Interactions Hypothesis (3, 4, 6). This association between mite domatia and temperate climes has also been found at smaller phylogenetic scales (23) and is recapitulated across published community surveys of domatia presence/absence. Indeed, a post hoc analysis using nine published studies with community surveys of mite domatia suggests a significantly higher proportions of species bear mite domatia in temperate compared to tropical zones (25 sites, $t = -2.347$, $df = 22.999$, $P\text{-value} = 0.028$, *SI Appendix, Fig. S7 and Table S4*). Thus, the most prominent hypothesis on investment in defense and mutualism with respect to latitude, which predicts more investment in equatorial zones, does not apply to

mite domatia, one of the most common plant-defense mutualism phenotypes. Rather, the type of plant–animal defense mutualism that is most common turns over from ant to mite as one moves poleward, reflecting distinct ecological associations in different regions rather than simply a pattern of overall higher investment in tropical environments.

Rather than tropicality per se, our analyses suggest that investment in mite domatia may instead be associated with climatic stress or mutualist availability. We found a negative correlation between mite domatia presence and rainfall, consistent with the hypothesis that mite domatia serve as adaptations for protecting beneficial mites from desiccation in dry environments (24). This pattern mirrors that found in elaiosome-bearing plants (i.e., plants that offer a food reward in exchange for dispersal by ants), which are more common in semiarid vs. tropical regions where plants are more water stressed and would likely benefit more from access to suitable germination sites afforded by ant dispersal (41). In contrast, two previously published intraspecific studies of mite domatia investment found positive associations between mite domatia size and precipitation (34, 35), indicating that environmental gradients in domatia investment within species may differ from broader patterns among species.

Among the most striking patterns we found were evolutionary correlations between mite domatia and other plant features, most notably woodiness and leaf phenology. The reason that mite domatia are so closely tied to woodiness remains a mystery. Interestingly, the very few exceptions of nonwoody plants with mite domatia in our dataset were nearly all found on domesticated crops (*SI Appendix, Table S2*), suggesting that the almost universal association between mite domatia and woodiness across natural systems can be broken by artificial selection and thus may not be due to a fundamental genetic or developmental constraint. Several adaptive explanations have been put forth but remain untested. O'Dowd and Willson (15) speculated that herbaceous species cannot provide mite populations with bark or other perennial shelters to serve as refugia during colder seasons (42, 43). While potentially important for temperate species, this mechanism does not explain the association between mite domatia and woody plants extending into tropical evergreen species, which do not experience winter. An alternative explanation draws from plant apparency theory, which posits that woody species are more likely to benefit from indirect defense traits than short-lived herbaceous species because it is easier for pests and bodyguards to discover woody species due to their long lifespans and large size (44). While these hypotheses are interesting, the strong association between mite domatia

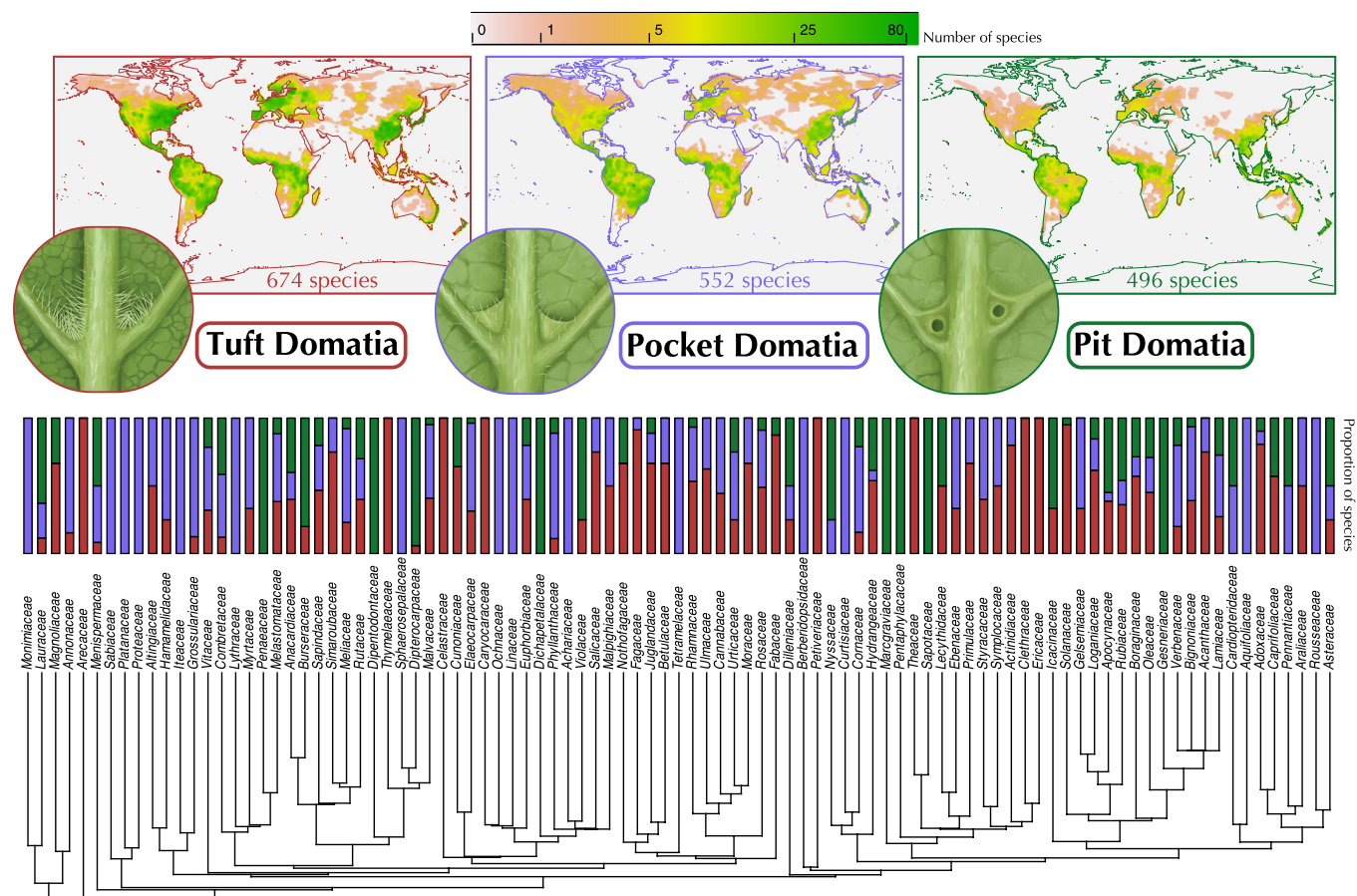


Fig. 3. Global patterns of occurrence of the three convergently evolved forms of domatia. Color shading in *Top* panels represents the number of species with that domatia type (tuft, pocket, or pit, from *Left* to *Right*), with greener colors denoting higher species richness. (*Bottom*) Percent abundance for the three morphological types of mite domatia for each domatia-bearing family. Domatia illustrations by John Megahan.

and woodiness represents an unanswered globally relevant botanical mystery, and experiments are needed to test the mechanism behind this striking evolutionary correlation.

Beyond an analysis of domatia presence and absence, our study is the first to rigorously evaluates the evolution of mite domatia morphotypes at a large-scale. We found a high degree of convergent evolution across the three main morphological forms of mite domatia (tufts, pockets, and pits). Twenty percent of families with domatia had representatives of all three mite domatia forms, with tuft form mite domatia associating with temperate, deciduous species and pit form domatia convergently associated with tropical regions in species with thicker, evergreen leaves. The mechanisms behind this pattern in mite domatia form remain an open question. However, given that convergence is a signature of adaptation, our findings strongly suggest domatia form is shaped by selection pressures that differ across regions. Future work focusing on quantifying the costs and benefits of different domatia forms across environmental contexts could tease apart whether selection pressures are biotic or abiotic. For example, pit domatia may render a leaf less vulnerable to colonization by pathogenic fungi than tuft domatia, as trichomes are known to provide surfaces amenable to fungal growth (45, 46). Different types of mite domatia could also be under selection by different mite taxa in different habitats. Reciprocal transplants among closely related plants that differ in their mite domatia morphology, as well as experiments that manipulate domatia form across habitats, are important next steps.

Future Directions and Conclusions. Understanding the evolutionary patterns of ecologically relevant traits is a central goal in biology. However, for many important traits, we lack the comprehensive understanding of their distribution needed to test hypotheses about their evolutionary causes and consequences across the tree of life. Until now, mite domatia—one of earth's most widespread mutualistic plant traits—was one of these phenotypes. Future work uncovering additional data on domatia presence, absence, and morphology will allow for the continued refinement of estimates of evolutionary rates of domatia gain and loss, as well as total numbers of domatia. Additional studies following up on this work using standardized methodologies (rather than consolidating disparate data sources) will be important for testing the generality and robustness of the patterns seen here across scales, geographies, and clades. Evaluating evolutionary gains and losses of domatia across well-sampled clades (e.g., genus-level studies), intraspecific studies that phenotype continuous variation within a species, and community-level studies that conduct standardized surveys across large environmental gradients are logical next steps to this work. Further, because our work revealed a clear pattern of evolutionary associations between mite domatia gain/loss and abiotic environmental variables, follow-up experiments focusing on the cost and benefits of domatia across environments will help illuminate the drivers of the phylogenetic correlations identified here.

Here, we have evaluated the broad phylogenetic and geographic distribution of mite domatia across all plants. Our results suggest that mite domatia occur in one out of every ten woody species

globally, having evolved a remarkable number of times convergently across the plant tree of life. Being more abundant in the temperate zone, the evolution and geographical distribution of mite domatia do not follow classical predictions; nonetheless our work suggests the notion that plant defense by mites transitions to plant defense by ants moving toward the tropics. This finding bears on the generality of theories for how and where species interactions evolve (5). In particular, given that particular geographic regions host distinct organismal communities and phenotypic forms, the biotic and abiotic attributes of these regions likely set the template for the differential evolution of adaptations related to species interactions. Defense mutualisms may be equally common in temperate and tropical environments, differing in their type rather than overall prevalence, representing distinct ecological associations and evolutionary histories.

Methods

World List of Mite Domatia-Bearing Species. We compiled a global list of plants with mite domatia using published reports and digital herbarium scans. We digitized data from two previously published lists which summarize accounts of mite domatia-bearing species reported from 1887 to 1990 (9, 11). To extend the list beyond previously published summaries, we conducted literature database surveys by querying Google Scholar (for articles published from 1990 through 2021) and the JSTOR Global Plants database (47) (all dates) with the search terms acarodomatia or the words "mite" and "domatia." The resulting data sources primarily included journal articles, books, theses and dissertations, and conference proceedings. We supplemented these sources by scoring dried plant specimens from herbaria for domatia phenotypes. To ensure phylogenetically broad sampling, we randomly phenotyped digitized herbarium scans in JSTOR Plants database (47) for the domatia from large woody angiosperm plant families (defined as families with >100 woody species according to Fitzjohn et al. (48) for which we had low representation in the literature search (<20 species in our literature search, 47 total families). We randomly sampled woody species from these families to bring the total number of sampled species to a minimum of 20 species for each angiosperm family that has greater than 100 woody species. For each scan, we scored up to three leaves per specimen. Species without leaves or with needle-like leaf morphology (e.g., Cactaceae, Proteaceae, etc.) were excluded.

Information extracted from all articles and scans included: taxon name down to the lowest possible taxonomic unit, mite domatia presence or absence, and mite domatia morphological category. Categories following Brouwer and Clifford, with tuft domatia being "dense cluster[s] of hairs"; pocket mite domatia consisting of a fold of "tissue connecting the diverging veins in the axil"; and pit mite domatia ranging "from shallow depressions to deep cavities with an opening at or near the [axil] center" (11). We were only able to classify mite domatia into morphotypes when photographs, diagrams, or detailed descriptions or references to other papers with these forms of evidence were available. When two or more reliable references described conflicting mite domatia types for a single species, we categorized the mite domatia type as "variable" (~2% of species in the dataset). Similarly, if the same species was described as having mite domatia in one reference and lacking mite domatia in another reference, or mite domatia were described as "sometimes," "often," or otherwise not always present, we treated the species as having mite domatia but made a note of the variability for future researchers (e.g., those interested in within species polymorphisms in domatia phenotypes, [Datasets S1 and S2](#)). Finally, some references—particularly taxonomic keys—indicated that all species within a clade either possessed or lacked mite domatia [e.g., all species in the genus *Perrottetia* (49)]. In these cases, we included all accepted species within the group as identified by the World Flora Online plant database (50). In accordance with Brouwer and Clifford (11), we considered more general plant structures that house mites but did not conform to the standard definition of mite domatia (e.g., rolled margins and midrib overhangs) as "pseudo mite domatia" and removed them from downstream analyses (listed in [SI Appendix, Table S6](#)).

We updated and standardized the taxonomic nomenclature of list entries using *WFO.match* and *WFO.one* functions in the R package *WorldFlora* (51),

which standardizes synonymous scientific names and resolves spelling errors. All "fuzzy matches" with a Levenshtein distance (i.e., "fuzzy.dist") values of 1 to 2 were accepted while values 3 and above were rejected and reverted to their originally assigned names. After updating species nomenclature, we collapsed all subspecies entries to the species for analysis and removed duplicate species entries. Cases where synonym identification and nomenclature updates revealed duplicate species with conflicting mite domatia information (presence/absence or morphology) were treated as conflicting accounts found in the literature as described above.

We acquired higher-level taxonomy information for each species using the *WorldFlora* package (51). The *WFO.match* function was used to assign species to families, and the *WFO.family* function to assign families to orders. Entries that were not matched to a family or order were entered manually. Entries that contained a genus name but no species name were removed from downstream analyses.

The Phylogenetic Distribution of Plants with Mite Domatia. We generated a species-level phylogeny of all the species in our dataset for downstream analyses using the *phylo.maker* "scenario 3" function in the *V.PhyloMaker* package (37) using the GBOTB extended mega-tree of vascular plants, which is based on clades built by Smith and Brown (52) and Zanne et al. (53). To visualize the phylogenetic distribution of mite domatia at the broad scale, we plotted domatia presence/absence on a trimmed phylogeny with one tip per family using the package *ape* (54). We summarized the broad distribution of domatia across taxonomic groups by calculating percentages of families and orders with mite domatia-bearing species using the most recent Angiosperm Phylogeny Group report (55).

We used Markov models of discrete trait evolution to estimate how often mite domatia were gained and lost over the evolutionary history of plants. We fit two models via maximum likelihood using *corHMM*: 1) a simple "all rates different" model with two states for lacking and bearing mite domatia and 2) a precursor model including an additional "hidden state" where lineages must first transition to an unobserved "precursor" state before being able to evolve mite domatia. We included the precursor model because 1) precursor models have been successful in modeling the evolution of other plant defense mutualism traits [i.e., EFNs; (56)], and 2) our data include both deep clades consisting entirely of species lacking domatia and recently diverged clades including both domatia-lacking and bearing species, strongly suggesting rates of domatia evolution vary among lineages. Unlike the simple model, the precursor model allows for such rate heterogeneity and thus exhibits a much better fit to our data ($\Delta AIC = 191.6$), so we report results based on the precursor model. We assumed the root state to be domatia-lacking (with no precursor), and we "down sampled" polytomies, retaining tips for each unique state (e.g., polytomies consisting of species entirely lacking/bearing mite domatia were collapsed to one tip, while polytomies with both states were collapsed to two tips with each state). We used *corHMM* to simulate stochastic character maps ("simmmaps") and infer ancestral states under each model as well as *phytools* to calculate the simulated number of transitions between states and visualized the inferred phylogenetic distribution of lineages with the precursor state (estimated to have evolved 160 to 210 times) using *shiftPlot*.

The Geographic Distribution of Plants with Mite Domatia. To describe contemporary geographic patterns of plants with mite domatia, we acquired occurrence data for species in our dataset using the GBIF (38, 39) using the R package *rgbif* (57). To ensure high data quality, we restricted the search to GBIF entries based on herbarium records and dropped dubious occurrences of coordinates that had uncertainty 100 km or greater, were outliers in terms of distances to other coordinates for the same species, were invalid, or were at GBIF headquarters, country capitals or centroids, or in oceans, using the *clean_coordinates* function in the R package *CoordinateCleaner* (58). We also removed any occurrences with a mismatch between reported coordinates and reported country of origin.

To evaluate the contemporary abiotic climatic conditions in which mite domatia-bearing species occur, we downloaded mean annual temperature and mean annual precipitation ([SI Appendix, Table S6](#)) data from WorldClim (59) using the *getData* function in the R package *sp* (60, 61) for all coordinates with available climate data. We acquired elevation data for all coordinates using the Google Maps Elevation API accessed using the *google_elevation* function in the *google-ewp* package (62). We replaced coastal negative elevation results with sea level

(0 elevation). Median values for latitude (absolute value), elevation, temperature, and precipitation were calculated for each species for use in downstream analyses.

We visualized the geographic distribution of species with mite domatia by reconstructing the global range of species in our dataset and creating rasters of the richness and phylogenetic diversity of species with mite domatia across the globe. We reconstructed concave hull ranges (for each species with more than three occurrences) and buffer ranges (for species with less than three occurrences) using buffer distances of 100,000 m using the *rangemap*, *hull*, and *raster* functions from the R package *rangemap* (63) and made raster diversity plots using the package *raster* (64).

Associations with Other Plant Traits. To test for relationships between mite domatia presence and other plant traits hypothesized to influence mite domatia evolution, we gathered data on three traits: leaf phenology; SLA; and plant woodiness. Data on woodiness and phenology were collated from the TRY Categorical Plant Traits Database (40), Fitzjohn et al. (48), Zanne et al. (53), and the TRY Leaf Phenology Database (65). SLA data were extracted from the Global Leaf Traits Database (65). We merged data from multiple sources by passing all angiosperm species names from these databases through the same *WorldFlora*-based pipeline used to clean the list of domatia-bearing species combining duplicate records for each species. Generally, species with conflicting or otherwise ambiguous records for categorical traits (e.g., "semievergreen") were classified as "variable." We combined SLA data by averaging measurements on the natural log scale after removing duplicate measurement records for a given species.

Statistical Analyses. To test whether the highly convergent evolutionary history of mite domatia reflected predictions from hypotheses about the drivers of mutualism, we conducted phylogenetic logistic regressions between mite domatia (presence/absence and morphology) and a) geographic factors (absolute value of latitude and elevation), b) climatic factors (mean annual temperature and total annual precipitation), and c) plant phenotypes (phenology and leaf thickness). We fit our data to phylogenetic logistic regression models (66) using the R package *phylolm* (67). We excluded known crop, ornamental, and known highly invasive species (68) from these analyses due to potential anthropogenic influence on their ranges and traits (Dataset S1). We ran separate univariate regressions for each predictor variable rather than a single multiple regression to avoid non-evenly distributed missing data across variables reducing the overall power of a combined analysis. We performed four regressions for each predictor variable: one "overall" analysis analyzing mite domatia absence and presence and three "morphotype" analyses analyzing the occurrence of each mite domatia morphotype (tuft, pocket, or pit) among all species with known morphotypes.

To assess significance, we simulated mite domatia occurrence and morphotypes using Markov models of discrete trait evolution and refit regressions to simulated data to assess how often our results would be observed if no relationship between mite domatia occurrence/morphology and climate/plant traits existed. This approach allows us to evaluate significance of our results while holding sampling across the phylogeny and geographic distributions of species constant, reducing potential biases due to species sampling. We estimated the maximum likelihood transition rates among all four states (i.e., mite domatia absence and all three morphotypes) under all rates different models using the R package *corHMM* (69). To avoid biased transition rate estimates due to polytomies (70), we collapsed polytomies into a single tip and treated these tips' states as ambiguous for all states exhibited by 1/k or more of tips in the original polytomy, where k is the total number of states observed across the tree. In coding these tips as ambiguous, we integrate across the uncertainty of tip states. Similarly, any species known to have mite domatia with unknown morphotype were treated as ambiguous for morphotype analyses. We used the R package *geiger* (71) to simulate mite domatia occurrence on the full phylogeny 1,000 times. Within polytomies, we resampled states such that the ranks of simulated state abundances within each polytomy for each simulation were largely conserved, while ensuring that distributions of state proportions matched that of the empirical data. We fit simulated data to phylogenetic logistic regression models. Any regressions that failed to converge (about 0 to 2% of regressions per predictor) were excluded from subsequent analyses. We used the *logspline* package (72) to estimate the probability that regressions fit to simulated data yield more extreme slope values than those from regressions fit to empirical data. Multiplying this probability by two, we obtain two-tailed *P*-values for observed relationships between mite domatia occurrence/

morphology and climate/plant traits. We applied the Benjamini–Hochberg correction to all 28 *P*-values (7 and 21 for the "overall" and "morphotype" regressions, respectively) to ensure a false discovery rate of ~5% (73). All data manipulation and analyses were performed using R version 4.1.1 (74).

Relationship with Ant Defense. To test for a trade-off between mite defense and ant defense across species and geography, we utilized data from Luo et al. (41), which collated and standardized data on plants with EFNs from Weber and Keeler (75) and plants with ant domatia (i.e., myrmecodomatia) from Chomicki and Renner (76). We created a combined dataset of plants scored as having mite defense (mite domatia) vs. ant defense (either EFNs or ant domatia) phenotypes. We assumed plants reported present for one trait but not the other as absent for the latter. To test for evolutionary correlations between mite and ant defense traits, we 1) fit hidden Markov models of binary character evolution using the R package *corHMM*, and 2) conducted phylogenetic logistic regressions using the same methods reported above (*Statistical Analyses*). For *corHMM* analyses, we allowed heterogeneity in the transition rates across the phylogeny to due hidden states, and the maximum likelihood of correlated and independent models was each computed using three random restarts, with the best-fitting model determined by AICc. To visualize geographic patterns in the relative abundance of plants with ant vs. mite defense traits, we calculated ranges of all species in the combined dataset as reported above (*The Geographic Distribution of Plants with Mite Domatia*) and plotted the number of species with mite domatia/the number of species with mite domatia plus the number of species with ant defense traits across a global raster, removing raster cells that had only singleton species reported. We tested for a relationship between the proportion of plants with mite vs. ant defense traits and latitude using an exponential regression using the *r* package *lm*. Specifically, we calculated the median latitude of each species using cleaned GBIF occurrences and regressed the proportion of species with any mutualistic trait (ant or mite) that possessed mite domatia against mean latitude, binning by three-degree latitudinal bands.

Estimating the Total True Number of Mite Domatia-Bearing Angiosperm Species. We derived an estimate of the total number of mite domatia-bearing woody angiosperm species (beyond those sampled in our list) using a taxonomically informed hierarchical Bayesian model. We focused on woody angiosperms (rather than all plants or all angiosperms) because nearly all known occurrences of mite domatia and thus the bulk of data available on which to base estimates are woody angiosperms (99.6%). This approach allowed us to avoid overestimating the total number of species with mite domatia across herbaceous groups and better estimate the number of domatia-bearing species in woody angiosperms. We jointly estimated the probability of having domatia alongside the probability of being herbaceous/woody for each angiosperm genus using the probabilistic programming language Stan (77). We assigned each species in our dataset to one of eight states: 0 *h*, 1 *h*, 0 *w*, 1 *w*, ? *h*, 0 ?, and 1 ?, where 0/1 indicates whether a species lacks or has domatia, *h/w* indicates whether a species is herbaceous or woody, and ? indicates when domatia/woodiness status is unknown (to accommodate uncertainty due to missing data). To counter potential biases in our data for literature sources reporting domatia presence over absence, we scored all species with known woodiness status in the TRY database but unknown domatia status as 0 *w*, incorporating the conservative assumption that species which are common enough to be scored for woodiness in TRY but do not have reports of domatia in the literature likely lack domatia. This decision was highly conservative, but it allowed us to calculate a minimum estimate for the number of woody species with leaf domatia given our known presence data, while avoiding any issues related to overestimation that could arise from missing absence data. We assumed counts of species in each state *i* within genus $J(n_{i,j})$ were multinomially distributed and estimated global intercepts with random effects for order, family, and genus to account for state probabilities being more similar among related taxa. We placed a normal prior with mean and SD 0 and 0.5, respectively, on the log-odds of being herbaceous (α_h) to reflect previous research suggesting slightly less than half of vascular plants are woody (48). We set conservative, normal priors with mean and SD -2 and 1.5, respectively on the log-odds of having domatia (α_r and α_s), and half-normal priors with SD 4 on all random effect SD parameters (β_{kl} , γ_{kl} , and δ_{kl}) to weakly regularize random effect estimates. For each model, posterior distributions were inferred by running four Hamiltonian Monte Carlo chains for 1,500 iterations, with the first 1,000 iterations as warmup and the last 500 retained as samples. Our analyses sampled the posterior distribution thoroughly, with \hat{R} 's < 1.01 and bulk and tail effective sample sizes >200 for over 99%

of parameters in each analysis. We used the inferred probabilities from the Bayesian hierarchical model to calculate estimates of the total number of domatia-bearing woody angiosperm species in each genus. Species in our dataset with partially observed states were first assigned to fully observed states by sampling binomial distributions with appropriate conditional probabilities (e.g., species in state 0? would be assigned to state 0h with probability $p_{0h}/(p_{0h} + p_{0w})$ and 0w otherwise). For genera with at least one observation, unobserved species were assigned to states by sampling from multinomial distributions with that genus' estimated probabilities. For genera with no observations, probabilities were drawn by sampling log-odds from normal distributions specified by estimated random effects (further ensuring that state probabilities, in the absence of additional information, will be more similar among closely related taxa). The R package *taxonlookup* (78) was used to identify the total number of accepted species in each genus. If a genus had more observed than accepted species (likely reflecting taxonomic disagreements among sources) we sampled and removed species without replacement to reduce the number of observed species to the number accepted. We repeated this procedure for each of the 2,000 posterior samples, yielding posterior distributions of counts of woody angiosperm species in each state.

1. G. Chomicki, M. Weber, A. Antonelli, J. Bascombe, E. T. Kiers, The impact of mutualisms on species richness. *Trends Ecol. Evol.* **34**, 698–711 (2019).
2. D. J. Futuyma, A. A. Agrawal, Macroevolution and the biological diversity of plants and herbivores. *Proc. Natl. Acad. Sci. U.S.A.* **106**, 18054–18061 (2009).
3. T. Dobzhansky, Evolution in the tropics. *Am. Sci.* **38**, 209–221 (1950).
4. J. L. Bronstein, R. Alarcón, M. Geber, The evolution of plant–insect mutualisms. *New Phytol.* **172**, 412–428 (2006).
5. D. W. Schemske, "Biotic interactions and speciation in the tropics" in *Speciation and Patterns of Diversity* (Cambridge University Press, 2009), pp. 219–239.
6. D. W. Schemske, G. G. Mittelbach, H. V. Cornell, J. M. Sobel, K. Roy, Is there a latitudinal gradient in the importance of biotic interactions? *Annu. Rev. Ecol. Syst.* **40**, 245–269 (2009).
7. R. C. Connor, The benefits of mutualism: A conceptual framework. *Biol. Rev.* **70**, 427–457 (1995).
8. M. D. Bertness, R. Callaway, Positive interactions in communities. *Trends Ecol. Evol.* **9**, 191–193 (1994).
9. A. N. Lundström, *Pflanzenbiologische Studien 2. Die Anpassungen der Pflanzen an Thiere* (Druck der Akademischen Buchdruckerei, Berlin, 1887).
10. S. A. Maccracken, I. M. Miller, C. C. Labandeira, Late Cretaceous domatia reveal the antiquity of plant–mite mutualisms in flowering plants. *Biol. Lett.* **15**, 20190657 (2019).
11. Y. M. Brouwer, H. T. Clifford, "An annotated list of domatia-bearing species" in *Notes from the Jodrell Laboratory* (Royal Botanic Gardens, Kew, 1990), pp. 1–33.
12. G. Q. Romero, W. W. Benson, Biotic interactions of mites, plants and leaf domatia. *Curr. Opin. Plant Biol.* **8**, 436–440 (2005).
13. G. English-Loeb, A. P. Norton, M. A. Walker, Behavioral and population consequences of acarodomatia in grapes on phytoseiid mites (Mesostigmata) and implications for plant breeding. *Entomol. Exp. Appl.* **104**, 307–319 (2002).
14. R. W. Pemberton, C. E. Turner, Occurrence of predatory and fungivorous mites in leaf domatia. *Am. J. Bot.* **76**, 105–112 (1989).
15. D. J. O'Dowd, M. F. Willson, Associations between mites and leaf domatia. *Trends Ecol. Evol.* **6**, 179–182 (1991).
16. D. E. Walter, D. J. O'Dowd, Leaf morphology and predators: Effect of leaf domatia on the abundance of predatory mites (Acari: Phytoseiidae). *Environ. Entomol.* **21**, 478–484 (1992).
17. R. Karban, G. English-Loeb, M. A. Walker, J. Thaler, Abundance of phytoseiid mites on *Vitis* species: Effects of leaf hairs, domatia, prey abundance and plant phylogeny. *Exp. Appl. Acarol.* **19**, 189–197 (1995).
18. J. A. M. Ferreira, B. Eshuis, A. Janssen, M. W. Sabelis, Domatia reduce larval cannibalism in predatory mites. *Ecol. Entomol.* **33**, 374–379 (2008).
19. A. A. Agrawal, R. Karban, Domatia mediate plantarthropod mutualism. *Nature* **387**, 562–563 (1997).
20. M. F. Willson, Foliar shelters for mites in the eastern deciduous forest. *Am. Midl. Nat.* **126**, 111–117 (1991).
21. D. J. O'Dowd, R. W. Pemberton, Leaf domatia in Korean plants: Floristics, frequency, and biogeography. *Vegetatio* **114**, 137–148 (1994).
22. D. J. O'Dowd, M. F. Willson, Leaf domatia and mites on Australasian plants: Ecological and evolutionary implications. *Biol. J. Linn. Soc.* **37**, 191–236 (1989).
23. M. G. Weber, W. L. Clement, M. J. Donoghue, A. A. Agrawal, Phylogenetic and experimental tests of interactions among mutualistic plant defense traits in *Viburnum* (Adoxaceae). *Am. Nat.* **180**, 450–463 (2012).
24. R. Grostal, D. J. O'Dowd, Plants, mites and mutualism: Leaf domatia and the abundance and reproduction of mites on *Viburnum tinus* (Caprifoliaceae). *Oecologia* **97**, 308–315 (1994).
25. A. P. Norton, G. English-Loeb, E. Belden, Host plant manipulation of natural enemies: Leaf domatia protect beneficial mites from insect predators. *Oecologia* **126**, 535–542 (2001).
26. G. A. Langellotto, R. F. Denno, Responses of invertebrate natural enemies to complex-structured habitats: A meta-analytical synthesis. *Oecologia* **139**, 1–10 (2004).
27. R. A. Schmidt, Leaf structures affect predatory mites (Acari: Phytoseiidae) and biological control: A review. *Exp. Appl. Acarol.* **62**, 1–17 (2014).
28. A. R. Wallace, *My Life* (Chapman & Hall, 1905).
29. Y. M. Brouwer, *Domatia: Morphology, Anatomy and Biological Significance* (The University of Queensland, 1985).
30. B. L. Bentley, Extrafloral nectaries and protection by pugnacious bodyguards. *Annu. Rev. Ecol. Syst.* **8**, 407–427 (1977).
31. M. Heil, D. McKey, Protective ant–plant interactions as model systems in ecological and evolutionary research. *Annu. Rev. Ecol. Syst.* **34**, 425–553 (2003).

Data, Materials, and Software Availability. Code for the analyses in this paper are deposited in Dryad (79).

ACKNOWLEDGMENTS. We are grateful to two reviewers whose comments substantially improved the manuscript. Thanks go to Susan Gordon for help with the literature search and Carina Baskett, Margaret Fleming, Carolyn Graham, Eric LoPresti, Dennis O'Dowd, Gideon Bradburd, James Boyko, Siyuyisiwe Situngu, Nathan Sanders, and members of the Weber Lab and the Michigan State University Plant Interactions Group for valuable feedback and help on the project. Thanks go to Laura Porturas for help translating list literature into text format and to the color blind simulator team at www.color-blindness.com for help checking our figures for accessibility. This research was funded in part by NSF grant numbers DEB-1831164 and DEB-2236747.

Author affiliations: ^aDepartment of Ecology and Evolutionary Biology, University of Michigan, Ann Arbor, MI 48103; ^bMichigan Nature Association, Okemos, MI 48864; ^cDepartment of Plant Biology, Program in Ecology, Evolution, and Behavior, Michigan State University, East Lansing, MI 48823; ^dDepartment of Evolution and Ecology, University of California, Davis, CA 95616; and ^eDepartment of Ecology and Evolutionary Biology, Cornell University, Ithaca, NY 14853

32. P. D. Coley, Effects of plant growth rate and leaf lifetime on the amount and type of anti-herbivore defense. *Oecologia* **74**, 531–536 (1988).
33. A. Pauw, *Mite-Plant Mutualism: Leaf Domatia of African Plants House Beneficial Mites* (University of Cape Town, 1992).
34. G. English-Loeb, A. Norton, Lack of trade-off between direct and indirect defence against grape powdery mildew in riverbank grape. *Ecol. Entomol.* **31**, 415–422 (2006).
35. L. A. Richards, P. D. Coley, Domatia morphology and mite occupancy of *Psychotria horizontalis* (Rubiaceae) across the Isthmus of Panama. *Arthropod. Plant Interact.* **6**, 129–136 (2012).
36. Y. Luo *et al.*, Climate and ant diversity explain the global distribution of ant–plant mutualisms. *Ecography* **2023**, e06841 (2023).
37. Y. Jin, H. Qian, V. Phylomaker, an R package that can generate very large phylogenies for vascular plants. *Ecography* **42**, 1353–1359 (2019).
38. GBIF.org, GBIF Homepage (2022). <https://www.gbif.org/>. Accessed 10 May 2022.
39. GBIF, Occurrence download. GBIF.org. <https://www.gbif.org/occurrence/download/0274722-210914110416597>. Accessed 10 May 2022.
40. J. Kattge *et al.*, TRY-Categorical traits dataset. Data from: TRY—a global database of plant traits. TRY File Archive. (2012). <https://www.try-db.org/TryWeb/Data.php>.
41. Y. Luo *et al.*, Climate and ant diversity explain the global distribution of ant–plant mutualisms. *Ecography* **2023**, e06841 (2023).
42. G. D. Broufas, D. S. Koveos, D. I. Georgatsis, Overwintering sites and winter mortality of *Euseius finlandicus* (Acari: Phytoseiidae) in a peach orchard in northern Greece. *Exp. Appl. Acarol.* **26**, 1–12 (2002).
43. M. Kawashima, C. Jung, Overwintering sites of the predacious mite *Neoseiulus californicus* (McGregor) (Acari: Phytoseiidae) in satsuma mandarin orchards on Jeju Island, Korea. *Appl. Entomol. Zool.* **45**, 191–199 (2010).
44. P. W. Price *et al.*, Interactions among three trophic levels: Influence of plants on interactions between insect herbivores and natural enemies. *Annu. Rev. Ecol. Syst.* **11**, 41–65 (1980).
45. J. Łażniewska, V. K. Macioszek, A. K. Kononowicz, Plant–fungus interface: The role of surface structures in plant resistance and susceptibility to pathogenic fungi. *Physiol. Mol. Plant Pathol.* **78**, 24–30 (2012).
46. K. W. Kim, Plant trichomes as microbial habitats and infection sites. *Eur. J. Plant Pathol.* **154**, 157–169 (2019).
47. JSTOR, JSTOR global plants database (2022). <https://plants.jstor.org/>. Accessed 11 May 2022.
48. R. G. FitzJohn *et al.*, How much of the world is woody? *J. Ecol.* **102**, 1266–1272 (2014).
49. K. Kubitzki, *Flowering Plants. Dicotyledons: Celastrales, Oxalidales, Rosales, Cornales, Ericales* (Springer Science & Business Media, 2004).
50. WFO, WFO (2022): World Flora online (2022). <http://www.worldfloraonline.org>. Accessed 11 May 2022.
51. R. Kindt, WorldFlora: An R package for exact and fuzzy matching of plant names against the World Flora Online taxonomic backbone data. *Appl. Plant Sci.* **8**, e11388 (2020).
52. S. A. Smith, J. W. Brown, Constructing a broadly inclusive seed plant phylogeny. *Am. J. Bot.* **105**, 302–314 (2018).
53. A. E. Zanne *et al.*, Three keys to the radiation of angiosperms into freezing environments. *Nature* **506**, 89–92 (2014).
54. E. Paradis, K. Schliep, ape 5.0: An environment for modern phylogenetics and evolutionary analyses in R. *Bioinformatics* **35**, 526–528 (2019).
55. Angiosperm Phylogeny Group *et al.*, An update of the Angiosperm Phylogeny Group classification for the orders and families of flowering plants: APG IV. *Bot. J. Linn. Soc.* **181**, 1–20 (2016).
56. B. Marazzi *et al.*, Locating evolutionary precursors on a phylogenetic tree. *Evol. Int. J. Org. Evol.* **66**, 3918–3930 (2012).
57. S. Chamberlain *et al.*, rgbif: Interface to the global biodiversity information facility API (R package 3.7.3, 2020). <https://CRAN.R-project.org/package=rgbif>. Accessed 10 May 2022.
58. A. Zizka *et al.*, "CoordinateCleaner: standardized cleaning of occurrence records from biological collection databases." *Methods Ecol. Evol.* **10**, 744–751 (2019).
59. S. E. Fick, R. J. Hijmans, WorldClim 2: New 1-km spatial resolution climate surfaces for global land areas. *Int. J. Climatol.* **37**, 4302–4315 (2017).
60. E. Pebesma, R. S. Bivand, Classes and methods for spatial data: The sp package. *R News* **5**, 9–13 (2005).
61. R. S. Bivand, E. J. Pebesma, V. Gómez-Rubio, E. J. Pebesma, *Applied Spatial Data Analysis with R* (Springer, 2008).

62. D. Cooley, googleway: Accesses Google Maps APIs to Retrieve Data and Plot Maps (R package version 2.0, 2017). <https://CRAN.R-project.org/package=googleway>. Accessed 10 May 2022.
63. M. E. Cobos, V. Barve, N. Barve, A. Jiménez-Valverde, C. Nuñez-Penichet. "Rangemap: An R package to explore species' geographic ranges". *Biodiv. Informatics*. **17**, 59–66 (2022).
64. R. Hijmans, raster: Geographic data analysis and modeling (R package version 3.6–30, 2024). <https://r-spatial.org/raster>. Accessed 4 September 2024.
65. J. Kattge *et al.*, TRY plant trait database—enhanced coverage and open access. *Glob. Change Biol.* **26**, 119–188 (2020).
66. A. R. Ives, T. Garland Jr., Phylogenetic logistic regression for binary dependent variables. *Syst. Biol.* **59**, 9–26 (2010).
67. L. S. T. Ho, C. Ane, A linear-time algorithm for Gaussian and non-Gaussian trait evolution models. *Syst. Biol.* **63**, 397–408 (2014).
68. Food and Agriculture Organization of the United Nations, *Programme for the World Census of Agriculture 2000* (FAO, Rome, 1995).
69. J. D. Boyko, J. M. Beaulieu, "Generalized hidden Markov models for phylogenetic comparative datasets." *Methods Ecol. Evol.* **12.3**, 468–478 (2021).
70. D. L. Rabosky, No substitute for real data: A cautionary note on the use of phylogenies from birth–death polytomy resolvers for downstream comparative analyses. *Evolution* **69**, 3207–3216 (2015).
71. M. W. Pennell *et al.*, geiger v2.0: An expanded suite of methods for fitting macroevolutionary models to phylogenetic trees. *Bioinformatics* **30**, 2216–2218 (2014).
72. C. Kooperberg, C. Moler, J. Dongarra, logspline: Routines for logspline density estimation (R Foundation for Statistical Computing, R package version 2.1.16, 2020). <https://cran.r-project.org/web/packages/logspline/index.html>. Accessed 1 June 2024.
73. Y. Benjamini, Y. Hochberg, Controlling the false discovery rate: A practical and powerful approach to multiple testing. *J. R. Stat. Soc. Ser. B Methodol.* **57**, 289–300 (1995).
74. R Core Team, R: A language and environment for statistical computing (R Foundation for Statistical Computing version 4.2.0., Vienna, Austria, 2022). <https://www.R-project.org/>. Accessed 1 May 2022.
75. M. G. Weber, K. H. Keeler, The phylogenetic distribution of extrafloral nectaries in plants. *Ann. Bot.* **111**, 1251–1261 (2013).
76. G. Chomicki, S. S. Renner, Phylogenetics and molecular clocks reveal the repeated evolution of ant-plants after the late Miocene in Africa and the early Miocene in Australasia and the Neotropics. *New Phytol.* **207**, 411–424 (2015).
77. B. Carpenter *et al.*, Stan: A probabilistic programming language. *J. Stat. Softw.* **76**, 1 (2017).
78. M. W. Pennell, R. G. FitzJohn, W. K. Cornwell, "A simple approach for maximizing the overlap of phylogenetic and comparative data." *Methods Ecol. Evol.* **7.6**, 751–758 (2016).
79. M. Weber, Data from "A global assessment of plant-mite mutualism and its ecological drivers." Dryad. <https://doi.org/10.5061/dryad.jq2bvq8jz>. Deposited 11 September 2024.